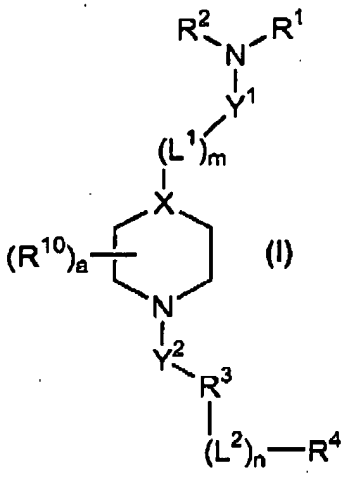
Listing of Claims

This listing of claims will replace all prior versions and listings of claims in the application.

Please cancel claim 11 without prejudice.

Please add new claims 19 and 20.

1. (Previously Presented) A compound of the formula (I)



(I)

wherein

a is an integer selected from 0 to 2;

R^{10} is selected from the group consisting of C_{1-6}alkyl, aryl, C_3-C_8cycloalkyl, aralkyl, heteroaryl, heteroaryl-C_{1-6}alkyl, heterocycloalkyl and heterocycloalky-C_{1-6}alkyl; wherein the aryl, cycloalkyl, aralkyl, heteroaryl or heterocycloalkyl group may be optionally substituted with one to four substituents independently selected from halogen, hydroxy, C_{1-6}alkyl, halogenated C_{1-6}alkyl, C_{1-6}alkoxy, halogenatedC_{1-6}alkoxy, nitro, cyano, amino, C_{1-4}alkylamino, di(C_{1-4}alkyl)amino, C_1.

$_6$alkylsulfonyl, C_{1-6}alkoxysulfonyl or halogenated C_{1-6}alkylsulfonyl;

X is selected from the group consisting of CH and C(C_1-C_6alkyl);

m is an integer selected from 0 and 1;

L^1 is selected from the group consisting of C_1-C_6alkyl;

Y^1 is selected from the group consisting of C(O) and C(S);

R^1 and R^2 are each independently selected from the group consisting of hydrogen, C_1-C_6alkyl, aryl, aralkyl, C_3-C_8cycloalkyl, C_3-C_8cycloalkyl-C_{1-6}alkyl, heteroaryl, heteroaryl-C_{1-6}alkyl, heterocycloalkyl and heterocycloalkyl-C_{1-6}alkyl; wherein the aryl, aralkyl, cycloalkyl, heteroaryl or heterocycloalkyl may be optionally substituted with one or more substituents independently selected from halogen, hydroxy, C_1-C_6alkyl, C_1-C_6alkoxy, halogenatedC_1-C_6alkyl, halogenatedC_1-C_6alkoxy, nitro, cyano, amino, C_1-C_4alkylamino, di(C_1-C_4alkyl)amino, heteroaryl or heterocycloalkyl;

alternatively, R^1 and R^2 may be taken together with the nitrogen atom to which they are bound to form a five to six membered monocyclic ring structure selected from the group consisting of pyrrolidinyl, piperidinyl, piperazinyl, morpholinyl and thiomorpholinyl;

Y^2 is selected from the group consisting of CH_2, C(O), C(S) and SO_2;

R^3 is selected from the group consisting of aryl and aralkyl; wherein the aryl or aralkyl may be optionally substituted with one of more substituents independently selected from halogen, hydroxy, C_1-C_6alkyl, C_1-C_6 alkoxy, halogenatedC_1-C_6alkyl, halogenatedC_1-C_6alkoxy, nitro, cyano, amino, C_1-C_4alkylamino, di(C_1-C_4alkyl)amino or $-(L^2)_m$-R^4;

n is an integer selected from 0 and 1;

L^2 is selected from the group consisting of C_1-C_8alkyl, C_2-C_8alkenyl, C_2-C_8alkynyl, C(O), C(S), SO_2 and $(A)_{0-1}—Q—(B)_{0-1}$;

where A and B are each independently selected from C_1-C_6alkyl, C_2-C_6alkenyl and C_2-C_6alkynyl;

where Q is selected from the group consisting of NR^5, O and S;

where R^5 is selected from the group consisting of hydrogen, C_1-C_6alkyl, aryl, aralkyl, C_{3-8}cycloalkyl, heteroaryl, heterocycloalkyl, C(O)-C_1-C_6alkyl, C(O)-aryl, C(O)-aralkyl, C(O)-heteroaryl, C(O)-heterocycloalkyl, SO_2-C_1-C_6alkyl, SO_2-aryl, SO_2-aralkyl, SO_2-heteroaryl, SO_2-heterocycloalkyl and $-CHR^6R^7$;

wherein the aryl, aralkyl, cycloalkyl, heteroaryl or heterocycloalkyl may be optionally substituted with one or more substituents independently selected from halogen, hydroxy, C_1-C_6alkyl, C_1-C_6 alkoxy, halogenatedC_1-C_6alkyl, halogenatedC_1-C_6alkoxy, nitro, cyano, amino, C_1-C_4alkylamino or di(C_1-C_4alkyl)amino;

where R^6 and R^7 are each independently selected from the group consisting of hydrogen, C_{1-6}alkyl, aryl, aralkyl, C_{3-8}cycloalkyl, heteroaryl, heterocycloalkyl, C(O)-C_{1-6}alkyl, C(O)aryl, C(O)-C_{3-8}cycloalkyl, C(O)-heteroaryl and C(O)-heterocycloalkyl; wherein the aryl, aralkyl, cycloalkyl, heteroaryl or heterocycloalkyl may be optionally substituted with one or more substituents independently selected from halogen, hydroxy, C_1-C_6alkyl, C_1-C_6 alkoxy, halogenatedC_1-C_6alkyl, halogenatedC_1-C_6alkoxy, nitro, cyano, amino, C_1-C_4alkylamino or di(C_1-C_4alkyl)amino;

R^4 is selected from the group consisting of aryl, aralkyl, C_3-C_8cycloalkyl, heteroaryl and heterocycloalkyl; wherein the aryl, aralkyl, cycloalkyl, heteroaryl or heterocycloalkyl may be optionally substituted with one or more substituents independently selected from halogen, hydroxy, C_1-C_6alkyl, C_1-C_6 alkoxy, halogenatedC_1-C_6alkyl, halogenatedC_1-C_6alkoxy, nitro, cyano, amino, C_1-C_4alkylamino or di(C_1-C_4alkyl)amino;

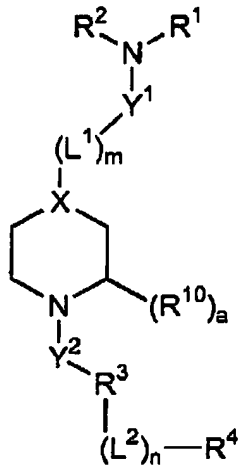
provided that when a is 0; X is CH; m is 1; L^1 is CH_2; R^3 is phenyl; n is 0; and R^4 is phenyl, wherein the phenyl group may be optionally substituted with one substituent selected from halogen, hydroxy, C_1-C_6alkyl, C_1-C_6 alkoxy, halogenatedC_1-C_6alkyl, halogenatedC_1-C_6alkoxy, nitro, cyano, amino, C_1-C_4alkylamino or di(C_1-C_4alkyl)amino, and wherein the R^4 group is bonded to the R^3 group in the para position;

then R^1 and R^2 are each independently selected from the group consisting of hydrogen, C_2-C_6alkyl, aryl, aralkyl, C_3-C_8cycloalkyl, C_3-C_8cycloalkyl-C_{1-6}alkyl, heteroaryl, heteroaryl-C_{1-6}alkyl, heterocycloalkyl and heterocycloalkyl-C_{1-6}alkyl; wherein the aryl, aralkyl, cycloalkyl, heteroaryl or heterocycloalkyl may be optionally substituted with one or more substituents independently selected from halogen, hydroxy, C_1-C_6alkyl, C_1-C_6alkoxy, halogenatedC_1-C_6alkyl, halogenatedC_1-C_6alkoxy, nitro, cyano, amino, C_1-C_4alkylamino, di(C_1-C_4alkyl)amino, heteroaryl or heterocycloalkyl;

alternatively, R^1 and R^2 may be taken together with the nitrogen atom to which they are bound to form a five to six membered monocyclic ring structure selected from the group consisting of pyrrolidinyl, piperidinyl, piperazinyl, morpholinyl and thiomorpholinyl;

and pharmaceutically acceptable salts thereof.

2. (Previously Presented) A compound as in Claim 1 of the formula



wherein

a is 0 to 1;

R^{10} is selected from the group consisting of C_1-C_4alkyl and aralkyl;

X is selected from the group consisting of CH and C(methyl);

m is an integer selected from 0 or 1;

L^1 is selected from the group consisting of C_1-C_4 alkyl;

Y^1 is C(O);

R^1 and R^2 are each independently selected from the group consisting of hydrogen, C_{1-4}alkyl, aryl, aralkyl, C_{3-8}cycloalkyl-C_1-C_4alkyl, heteroaryl and heterocycloalkyl; wherein the aryl, aralkyl or heteroaryl may be optionally substituted with one to two substituents independently selected from halogen, hydroxy,

C_1-C_4alkyl, C_1-C_4alkoxy, trifluoromethyl, trifluoromethoxy, C_1-C_4alkylamino, di(C_1-C_4alkyl)amino or heterocycloalkyl;

alternatively, R^1 and R^2 may be taken together with the nitrogen atom to which they are bound to form a five to six membered monocyclic ring structure selected from the group consisting of pyrrolidinyl, piperidinyl, piperazinyl, morpholinyl and thiomorpholinyl; Y^2 is C(O);

R^3 is selected from the group consisting of aryl; wherein the aryl may be optionally substituted with one to two substituents independently selected from C_1-C_4alkyl, trifluoromethyl or $-(L^2)_n-R^4$;

n is an integer selected from 0 or 1;

L^2 is selected from the group consisting of C_1-C_6alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl and $(A)_{0-1}—Q—(B)_{0-1}$;

where A and B are each independently selected from C_1-C_4alkyl;

where Q is selected from the group consisting of NR^5, O and S;

where R^5 is selected from the group consisting of hydrogen, C_1-C_4alkyl, C(O)-C_1-C_6alkyl, C(O)-aryl, C(O)-aralkyl, C(O)-heteroaryl, C(O)-heterocycloalkyl and $-CHR^6R^7$; wherein the aryl, aralkyl, cycloalkyl, heteroaryl or heterocycloalkyl may be optionally substituted with one to two substituents independently selected from halogen, C_1-C_4alkyl, C_1-C_4alkoxy, trifluoromethyl, trifluoromethoxy, nitro, cyano, amino, C_1-C_4alkylamino or di(C_1-C_4alkyl)amino;

where R^6 and R^7 are each independently selected from the group consisting of hydrogen, C_{1-4}alkyl, aryl, aralkyl, C_{3-8}cycloalkyl, heteroaryl, heterocycloalkyl, C(O)-C_{1-6}alkyl,

C(O)aryl, C(O)-C_{3-8}cycloalkyl, C(O)-heteroaryl and C(O)-heterocycloalkyl; wherein the aryl, aralkyl, cycloalkyl, heteroaryl or heterocycloalkyl may be optionally substituted with one to two substituents independently selected from halogen, hydroxy, C_1-C_4alkyl, C_1-C_4alkoxy, trifluoromethyl, trifluoromethoxy, nitro, cyano, amino, C_1-C_4alkylamino or di(C_1-C_4alkyl)amino;

R^4 is selected from the group consisting of aryl, heteroaryl and heterocycloalkyl; wherein the aryl group may be optionally substituted with one to two substituents independently selected from halogen, hydroxy, C_1-C_4alkyl, C_{1-4}alkoxy, trifluoromethyl or amino;

provided that when a is 0; X is CH; m is 1; L^1 is CH_2; R^3 is phenyl; n is 0; and R^4 is phenyl, wherein the phenyl group may be optionally substituted with one substituent selected from halogen, hydroxy, C_1-C_4alkyl, C_1-C_4alkoxy, trifluoromethyl or amino, and wherein the R^4 group is bonded to the R^3 group in the para position;

then R^1 and R^2 are each independently selected from the group consisting of hydrogen, C_{2-4}alkyl, aryl, aralkyl, C_{3-8}cycloalkyl-C_1-C_4alkyl, heteroaryl and heterocycloalkyl; wherein the aryl, aralkyl or heteroaryl may be optionally substituted with one to two substituents independently selected from halogen, hydroxy, C_1-C_4alkyl, C_1-C_4alkoxy, trifluoromethyl, trifluoromethoxy, C_1-C_4alkylamino, di(C_1-C_4alkyl)amino or heterocycloalkyl;

alternatively, R^1 and R^2 may be taken together with the nitrogen atom to which they are bound to form a five to six membered monocyclic ring structure selected from the group

consisting of pyrrolidinyl, piperidinyl, piperazinyl, morpholinyl and thiomorpholinyl;

and pharmaceutically acceptable salts thereof.

3. (Previously Presented) A compound as in Claim 2 wherein

X is CH;

m is 1;

R^1 is selected from the group consisting of hydrogen and C_{1-4}alkyl;

R^2 is selected from the group consisting of C_{1-4}alkyl, aryl, aralkyl, C_{3-8}cycloalkyl-C_{1-4}alkyl and heteroaryl; wherein the aryl or aralkyl may be optionally substituted with one to two substituents independently selected from halogen, hydroxy, C_1-C_4alkyl, C_1-C_4alkoxy, trifluoromethyl, trifluoromethoxy, di(C_1-C_4alkyl)amino or heterocycloalkyl;

alternatively, R^1 and R^2 may be taken together with the nitrogen atom to which they are bound to form a five to six membered monocyclic ring structure selected from the group consisting of pyrrolidinyl, piperidinyl and morpholinyl;

R^3 is selected from the group consisting of aryl; wherein the aryl may be optionally substituted with a substituent selected from C_1-C_4alkyl or trifluoromethyl;

L^2 is selected from the group consisting of C_1-C_4alkyl, C_2-C_6alkenyl, C_2-C_6alkynyl, NH-C_{1-4}alkyl, C_{1-4}alkyl-N(C_{1-4}alkyl)-C_{1-4}alkyl and C_{1-4}alkyl-N(C(O)C_{1-4}alkyl)-C_{1-4}alkyl;

provided that when a is 0; X is CH; L^1 is CH_2; R^3 is phenyl; n is 0; and R^4 is phenyl, wherein the phenyl group may be optionally substituted with one substituent selected from

halogen, hydroxy, C_1-C_4alkyl, C_1-C_4alkoxy, trifluoromethyl or amino, and wherein the R^4 group is bonded to the R^3 group in the para position;

then R^1 is selected from the group consisting of hydrogen and C_{2-4}alkyl;

R^2 is selected from the group consisting of C_{2-4}alkyl, aryl, aralkyl, C_{3-8}cycloalkyl-C_{1-4}alkyl and heteroaryl; wherein the aryl or aralkyl may be optionally substituted with one to two substituents independently selected from halogen, hydroxy, C_1-C_4alkyl, C_1-C_4alkoxy, trifluoromethyl, trifluoromethoxy, di(C_1-C_4alkyl)amino or heterocycloalkyl;

alternatively, R^1 and R^2 are taken together with the nitrogen atom to which they are bound to form a five to six membered monocyclic ring structure selected from the group consisting of pyrrolidinyl, piperidinyl and morpholinyl;

and pharmaceutically acceptable salts thereof.

4. (Previously Presented) A compound as in Claim 3 wherein

R^{10} is selected from the group consisting of methyl and benzyl;

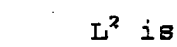
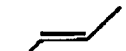
L^1 is selected from the group consisting of CH_2 and CH_2CH_2;

R^2 is selected from the group consisting of -CH_2-(3-trifluoromethylphenyl), -CH_2-cyclohexyl, -CH_2-(3,5-dimethoxyphenyl), -CH_2-(4-trifluoromethylphenyl), -CH_2-(3,5-ditrifluoromethylphenyl), 3-trifluoromethoxyphenyl, -CH_2-(4-dimethylaminophenyl), phenyl, benzyl, 2-fluorophenyl, 4-fluorophenyl, 2,4-difluorophenyl, 2,6-difluorophenyl, 4-hydroxyphenyl, 4-dimethylamino-phenyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, 4-pyridyl-methyl, 4-morpholinyl-phenyl, 4-piperidinyl-

phenyl, methyl, isopropyl, 4-methoxyphenyl, 4-trifluoromethylphenyl, 2-pyrimidinyl, 4-pyrimidinyl,5-quinolinyl, 6-quinolinyl, and 8-quinolinyl;

alternatively, R^1 and R^2 are taken together with the nitrogen atom to which they are bound to form a five to six membered monocyclic ring structure selected from the group consisting of pyrrolidinyl, piperidinyl and morpholinyl;

R^3 is selected from the group consisting of phenyl, methylphenyl and trifluoromethylphenyl;

L^2 is selected from the group consisting of 2-  , 3-, 4-, 5-, 2-  , 3-  , 2-  , 3-  , 4-  , 2-CH_2CH_2, 3-CH_2-CH_2, 4-CH_2-CH_2, NH-CH_2, CH_2-N(CH_3)-CH_2, CH_2-N(CH_3)-CH_2CH_2, CH_2-N(C(O)CH_3)-CH_2 and CH_2-N(C(O)CH_3)-CH_2CH_2;

R^4 is selected from the group consisting of phenyl, 1-naphthyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, 3-hydroxyphenyl, 2-methylphenyl, 3-aminophenyl, 4-methoxyphenyl, 4-chlorophenyl, 2-thienyl, 3-thienyl, 3,5-di(trifluoromethyl)-phenyl, 1-imidazolyl, 2-benzimidazolyl, 1-pyrrolidinyl, 2-furyl and 2-tetrahydrofuryl;

provided that when a is 0; X is CH; L^1 is CH_2; R^3 is phenyl; n is 0; and R^4 is phenyl, 4-chlorophenyl, 3-hydroxyphenyl, 2-methylphenyl, 4-methoxyphenyl or 3-aminophenyl; and wherein the R^4 group is bonded to the R^3 group in the para position;

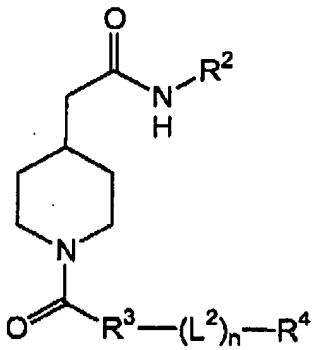
then R^1 is selected from the group consisting of hydrogen and C_{2-4}alkyl;

R^2 is selected from the group consisting of $-CH_2$-(3-trifluoromethylphenyl), $-CH_2$-cyclohexyl, $-CH_2$-(3,5-dimethoxyphenyl), $-CH_2$-(4-trifluoromethylphenyl), $-CH_2$-(3,5-ditrifluoromethylphenyl), 3-trifluoromethoxyphenyl, $-CH_2$-(4-dimethylaminophenyl), phenyl, benzyl, 2-fluorophenyl, 4-fluorophenyl, 2,4-difluorophenyl, 2,6-difluorophenyl, 4-hydroxyphenyl, 4-dimethylamino-phenyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, 4-pyridyl-methyl, 4-morpholinyl-phenyl, 4-piperidinyl-phenyl, isopropyl, 4-methoxyphenyl, 4-trifluoromethylphenyl, 2-pyrimidinyl, 4-pyrimidinyl,5-quinolinyl, 6-quinolinyl, and 8-quinolinyl;

alternatively, R^1 and R^2 are taken together with the nitrogen atom to which they are bound to form a five to six membered monocyclic ring structure selected from the group consisting of pyrrolidinyl, piperidinyl and morpholinyl;

and pharmaceutically acceptable salts thereof.

5. (Original) A compound as in Claim 4 of the formula



wherein

R^2 is selected from the group consisting of $-CH_2$-(3-trifluoromethylphenyl), $-CH_2$-cyclohexyl, $-CH_2$-(3,5-dimethoxyphenyl), $-CH_2$-(4-trifluoromethylphenyl), $-CH_2$-(3,5-

ditrifluoromethylphenyl), -CH_2-(4-dimethylaminophenyl), phenyl, 2-fluorophenyl, 4-fluorophenyl, 2,4-difluorophenyl, 2,6-difluorophenyl, 3-trifluoromethylphenyl, 4-trifluoromethylphenyl, 4-hydroxyphenyl, 4-methoxyphenyl, benzyl, 3-pyridyl, 4-pyridyl, 2-pyrimidinyl, 4-pyrimidinyl, 5-quinolinyl, 6-quinolinyl, 8-quinolinyl, 4-(dimethylamino)-phenyl, 4-morpholinyl-phenyl, 4-pyridyl-methyl, and 4-piperidinyl-phenyl;

L^2 is selected from the group consisting of 2-  , 3-, 4-, 5-, 2-  , 3-  , 4-  , 2-  , 3-  , 2-CH_2CH_2, 3-CH_2-CH_2, 4-CH_2-CH_2, NH-CH_2, 4-(CH_2-N(CH_3)-CH_2), 4-(CH_2-N(CH_3)-CH_2CH_2), 4-(CH_2-N(C(O)CH_3)-CH_2) and 4-(CH_2-N(C(O)CH_3)-CH_2);

R^4 is selected from the group consisting of phenyl, 3-phenyl; 5-phenyl, 4-chlorophenyl, 3-hydroxyphenyl, 3-(2-methylphenyl), 3-(3-aminophenyl), 2-pyridyl, 3-pyridyl, 3-(3-pyridyl), 4-pyridyl, 3-(3-thienyl), 3,5-di(trifluoromethyl)phenyl, 1-pyrrolidinyl, 2-furyl, 1-naphthyl, 2-thienyl, 1-imidazolyl, 2-benzimidazolyl and 2-tetrahydrofuryl;

and pharmaceutically acceptable salts thereof.

6. (Canceled)

7. (Original) A compound as in Claim 4 selected from the group consisting of

N-phenyl-1-[3-(2-pyridinylethynyl)benzoyl]-4-piperidineacetamide;

N-(2,4-difluorophenyl)-1-[3-(2-pyridinylethynyl)benzoyl]-4-piperidineacetamide;

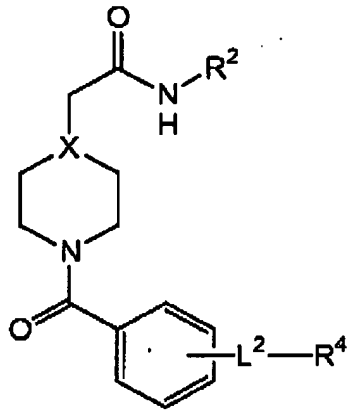
N-phenyl-4-[2-[(*E*)-2-(2-pyridinyl)ethenyl]benzoyl]-1-piperazineacetamide;

N-phenyl-4-[3-(2-pyridinylethynyl)benzoyl]-1-piperazineacetamide;

N-(4-hydroxyphenyl)-1-[3-(2-pyridinylethynyl)benzoyl]-4-piperidineacetamide;

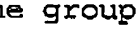
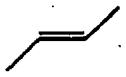
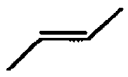
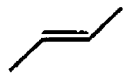
and pharmaceutically acceptable salts thereof.

8. (Previously Presented) A compound as in Claim 4 wherein of the formula



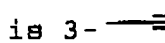
X is CH;

R^2 is selected from the group consisting of phenyl, 4-hydroxyphenyl, 2-fluorophenyl, 4-fluorophenyl, and 2,4-difluorophenyl;

L^2 is selected from the group consisting of 3-  , 4-  , 2-  , 3-  , 4-  , 4-(CH_2-N(CH_3)-CH_2CH_2), 4-(CH_2-N(CH_3)-CH_2) and 3-NH-CH_2;

R^4 is selected from the group consisting of 2-pyridyl, 4-pyridyl, 4-pyrrolidinyl, 2-furyl, 1-naphthyl and 3,5-di(trifluoromethyl)phenyl;

and pharmaceutically acceptable salts thereof.

9. (Original) A compound as in Claim 8 wherein X is CH; R^2 is phenyl; L^2 is 3-  ; R^4 is 2-pyridyl and pharmaceutically acceptable salts thereof.

10. (Original) A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a compound of Claim 1.

11. (Canceled).

12. (Original) A process for making a pharmaceutical composition comprising mixing a compound of Claim 1 and a pharmaceutically acceptable carrier.

13. (Original) A method of treating a nervous system disorder in a subject in need thereof comprising administering to the subject a therapeutically effective amount of the compound of Claim 1.

14. (Previously Presented) The method of Claim 13, wherein the nervous system disorder is selected from the group consisting of depression, dementia, schizophrenia, bipolar disorders, anxiety, emesis, acute pain, neuropathic pain, itching, migraine and movement disorders.

15. (Original) A method of treating nervous system a disorder in a subject in need thereof comprising administering to the subject a therapeutically effective amount of the composition of Claim 10.

16. (Original) A method of treating a nervous system disorder selected from the group consisting of depression and anxiety in a subject in need thereof comprising administering to the subject a therapeutically effective amount of the compound of Claim 1.

17. (Original) A method of treating a nervous system disorder selected from the group consisting of depression and anxiety in a subject in need thereof comprising administering to the subject a therapeutically effective amount of the pharmaceutical composition of Claim 10.

18. (Original) A method of treating a nervous system disorder selected from the group consisting of depression and anxiety in a subject in need thereof comprising administering to the subject a therapeutically effective amount of the compound of Claim 9.

19. (New) The compound of claim 1 wherein R^4 is selected from the group consisting of phenyl, 2-pyridyl, 3-pyridyl, 4-pyridyl, 3-hydroxyphenyl, 2-methylphenyl, 3-aminophenyl, 3-thienyl, 3,5-di(trifluoromethyl)phenyl, 4-methoxyphenyl, 4-chlorophenyl, 2-thienyl, 2-furyl, 1-pyrrolidinyl,1-imidazolyl, 2-benzimidazolyl, naphthyl and 2-tetrahydrofuryl.

20. (New) The compound of claim 1 wherein R^4 is selected from the group consisting of phenyl, 3-phenyl; 5-phenyl, 4-chlorophenyl, 3-hydroxyphenyl, 3-(2-methylphenyl), 3-(3-aminophenyl), 2-pyridyl, 3-pyridyl, 3-(3-pyridyl), 4-pyridyl, 3-(3-thienyl), 3,5-di(trifluoromethyl)phenyl, 1-pyrrolidinyl, 2-furyl, 1-naphthyl, 2-thienyl, 1-imidazolyl, 2-benzimidazolyl and 2-tetrahydrofuryl.